

JC625 U.S. PTO
09/523532
03/10/00

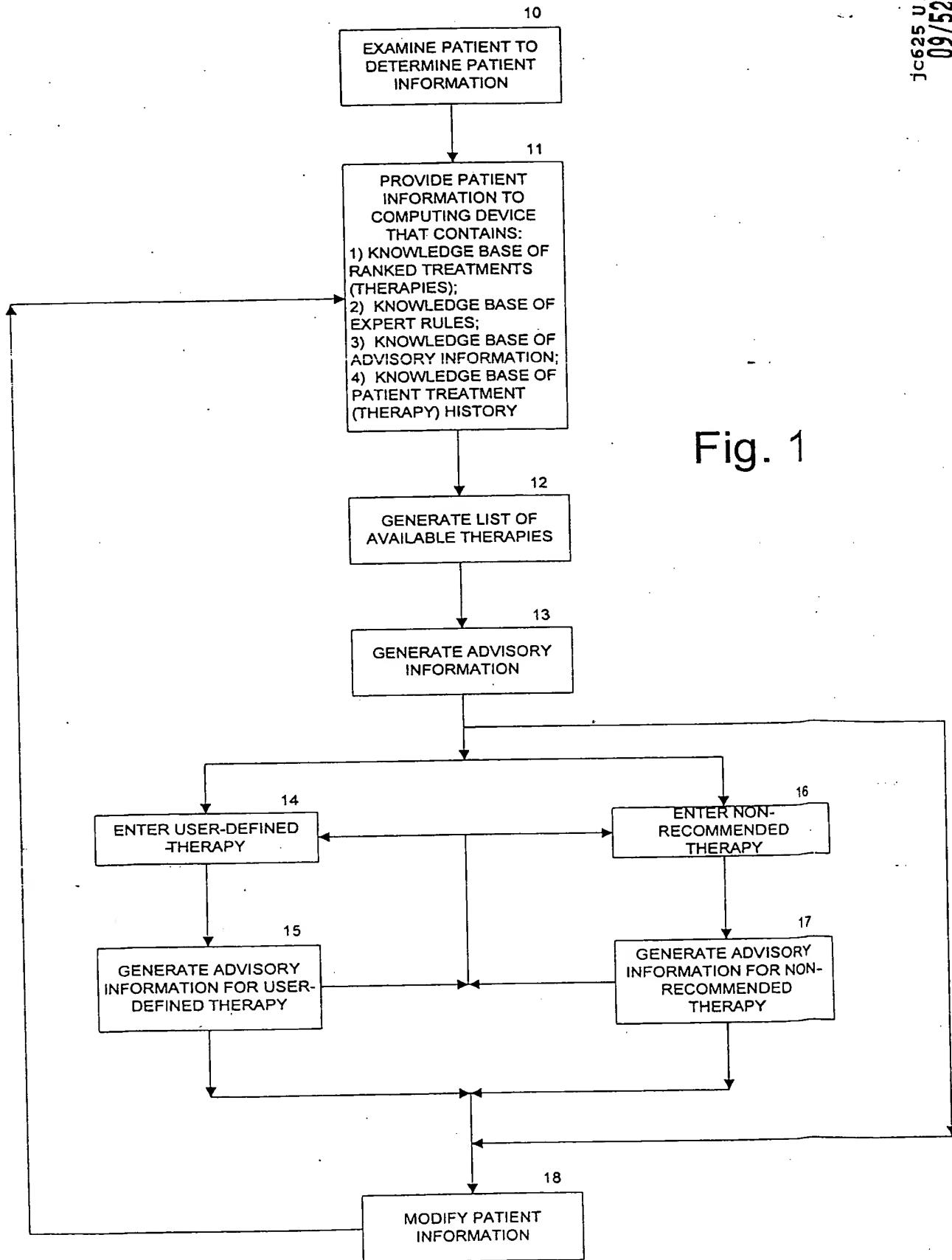


Fig. 1

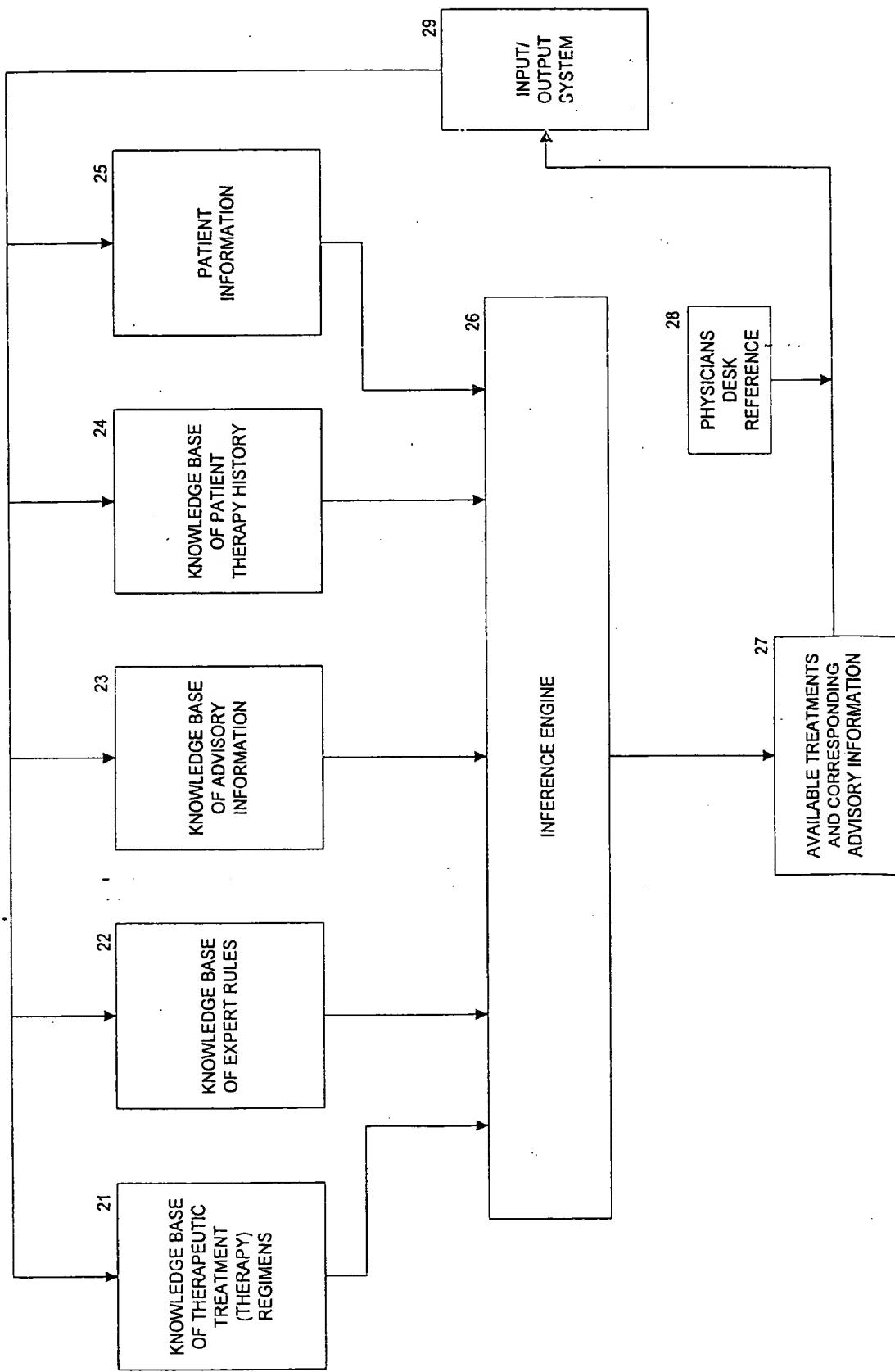


Fig. 2

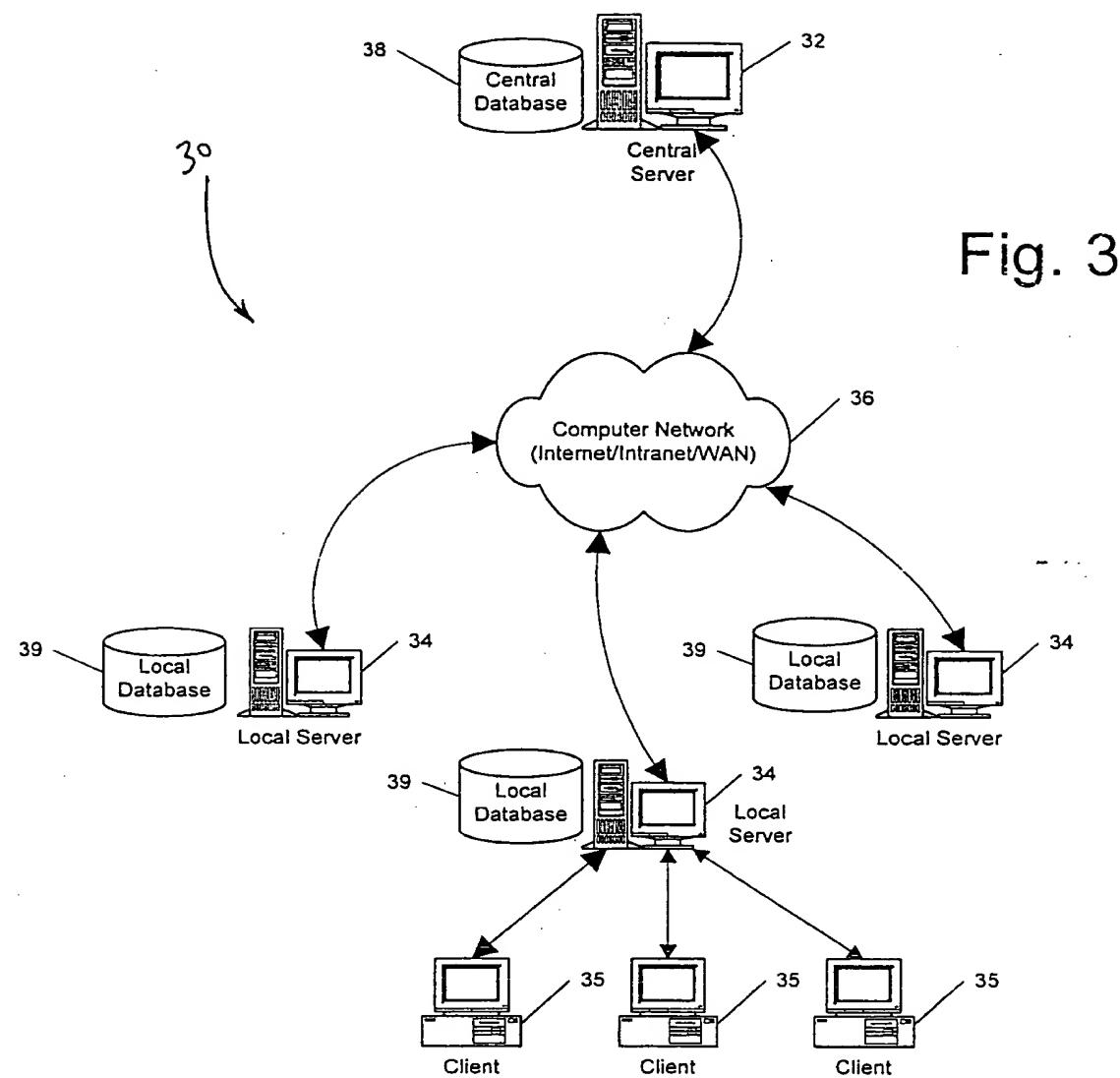


Fig. 3

50a 60a 70a 51 52 53

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Medical History Chart / Therapy Evaluation

General

POLYMER LETTERS EDITION

IFMS Number

Gender Copy Date

End Viral Load

Specimen No. _____ Date _____ Value _____

人間の精神は、常に進歩するものである。

Specimen Date _____

HISTOIRE DE LA
LIVRE

Pheno type

ARVANITOU

Intoleranță

Неманъ

Specimen Date _____ Value (g/dL) _____

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Pancratius
Neurophysiol

Specimen Date: _____

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Specimen Date: AST/SG01/11/11

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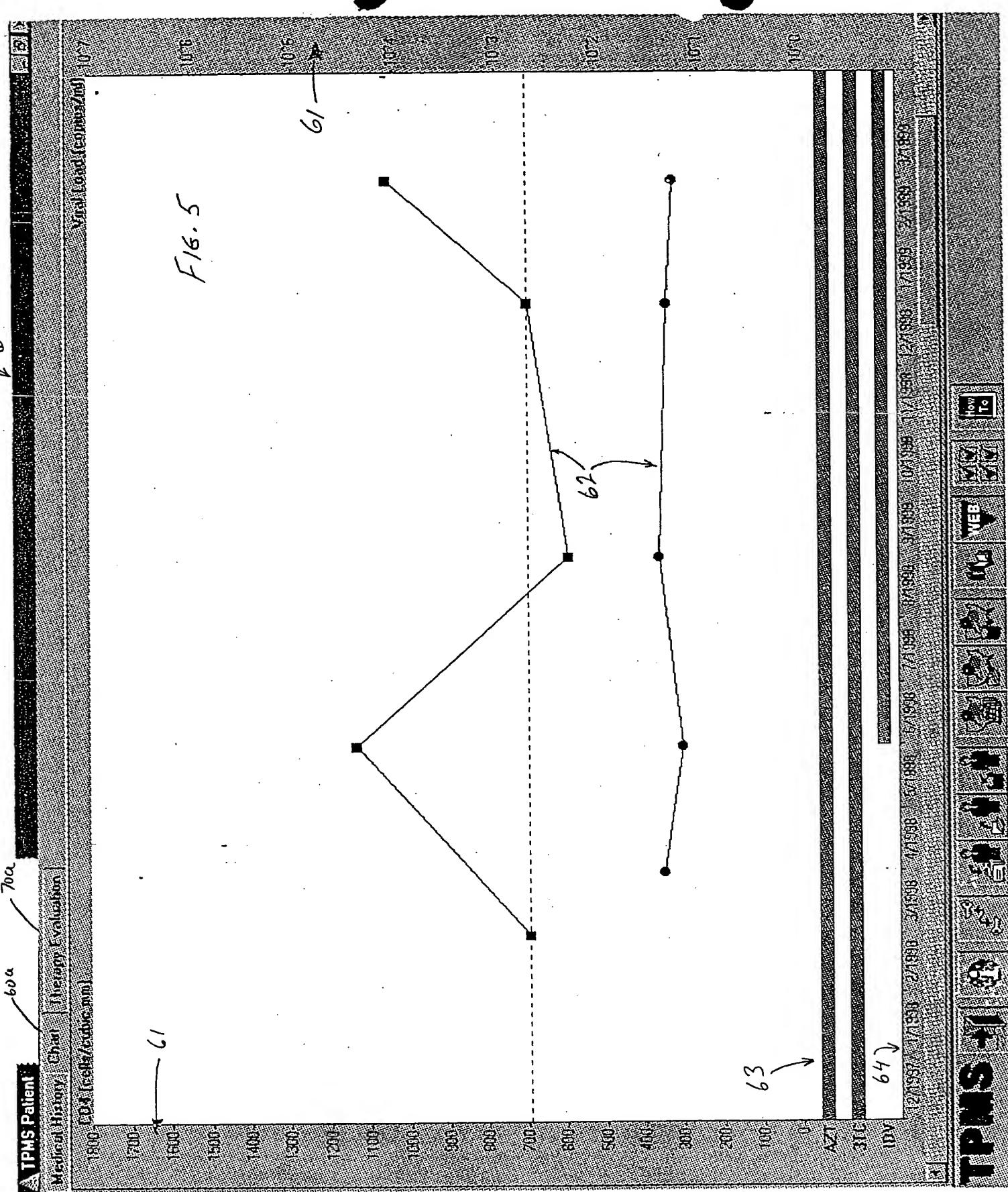
THE ECONOMIC FORECASTS OF THE INSTITUTE FOR INTERNATIONAL TRADE 11

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F16. 4



F16. 7

Icon	Meaning
⊕	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box.
⊕	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box. The book indicates that therapy has been studied and a reference is available to review.
△	Indicates a yellow alert. There is important information about this therapy that must be reviewed.
△	Indicates a yellow alert. There is important information about this therapy that must be reviewed. The book indicates that therapy has been studied and a reference is available to review.
!	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered.
!	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered. The book indicates that therapy has been studied and a reference is available to review.
✗	Indicates the therapy is not recommended.

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IPMS Patient

Medical History	Chemotherapy/Evaluation
Therapy Being Evaluated	

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Read the following Red Drug Contra-Indication Alerts for this therapy:

- **Drug Interaction Alert:** Patient is currently taking zolpidem, co-administration of Norvir (Ritonavir/RTV) with certain non-sedating antihistamines, sedative hypnotics, or oral contraceptives may result in potentially serious and/or life-threatening adverse events due to possible effects of Norvir (Ritonavir/RTV) on the hepatic metabolism of certain drugs. Norvir (Ritonavir/RTV) can produce large increases in plasma concentrations of certain highly metabolized drugs. Norvir (Ritonavir/RTV) should not be coadministered with alprazolam, amiodarone, astemizole, bisoprolol, buproprion, citalopram, clorazepate, diazepam, doxepin, fluoxetine, fexofenadine, fexofenadine, triazolam or zolpidem. Patient is taking zolpidem and in order to use this therapy, that drug should be replaced with a non-contraindicated substitute. CmDQ, Commentary 25

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Dosages

- Ritonavir 200mg q12h (2 pills/day), \$2,56/day*
- Valtrex 125mg q12h (4 pills/day), \$1,22/day*
- Invacare 400mg q12h; taken within 2 hours after a full meal (4 pills/day, \$3,17/day)
- Norvir 400mg q12h (3 pills/day, \$14.84/day)

(* indicates adjusted dosage)

30 Dosage Adjustments: The following dosage adjustments messages apply to this therapy:

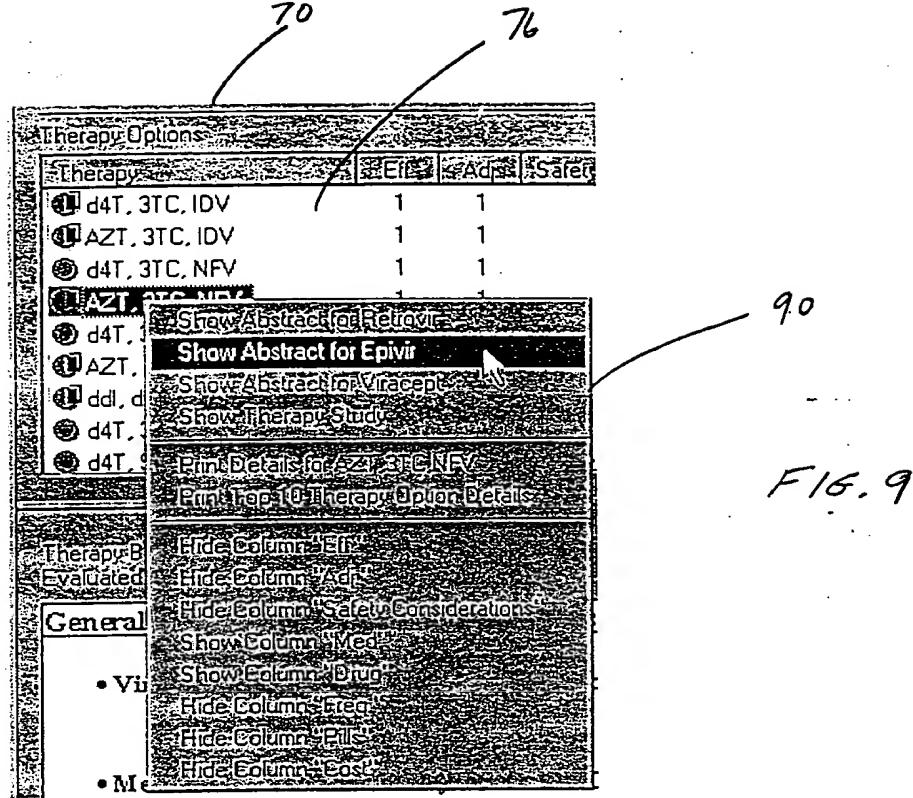
- **Dosage Notice:** This therapy contains both sequinavir and rilonavir. When rilonavir and sequinavir are used together the dosage of each drug is reduced by 1/2. The dosage for these drugs has been set accordingly. DxD-C and, Commentary 26

Invacare (quinavir/SQV): The following Warnings and Adverse effects apply to Invacare (quinavir/SQV):

- **Drug Interactions Information:** Compounds that are substrates of CYP3A4 (e.g., calcium channel blockers, clindamycin, dapsone, quinidine, in soltan) may have elevated plasma concentrations when coadministered with Invacare (quinavir/SQV); therefore, patient should be monitored for toxicities associated with such drugs when taking Invacare (quinavir/SQV). CmDQ, Commentary 21

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F16.8



TPMS F

Hematological Data

Therapy Evaluation

General

Patient Id	<input type="text"/> demo1	Birth Date	<input type="text"/> 1/1/1960	TPMS Number	<input type="text"/>
Physician	<input type="text"/>	Gender	<input type="text"/> Male	First	<input type="checkbox"/>

CD4 and Viral Load

CD4 (cells/cubic mm)	Specimen Date	Date	Specimen Description	Specimen Value
100	<input type="text"/> 3/1/1999	<input type="text"/> 3/1/1999	<input type="text"/> 320	<input type="text"/> 340
1000	<input type="text"/> 3/1/1999	<input type="text"/> 3/1/1999	<input type="text"/> 12000	<input type="text"/> C/mL
Previous Viral Load	<input type="text"/> 1/1/1999	<input type="text"/> 1/1/1999	<input type="text"/> 500	<input type="text"/> 1000

HIV Genotype

Phenotype	<input type="text"/>	<input type="text"/>
Allergy/Hyper	<input type="text"/>	<input type="text"/>
Tolerance	<input type="text"/>	<input type="text"/>
	<input type="text"/>	<input type="text"/>

Hemoglobin

Specimen Date	Value	Date	Value
<input type="text"/> 3/1/1999	<input type="text"/> 12.00	<input type="text"/> 3/1/1999	<input type="text"/> No

Neutrophils

Specimen Date	Value	Date	Value
<input type="text"/> 3/1/1999	<input type="text"/> 1500	<input type="text"/> 3/1/1999	<input type="text"/> No

Renal Function

Specimen Date	Value	Date	Specimen Date	Specimen Value	Explanations
<input type="text"/> 3/1/1999	<input type="text"/> 49	<input type="text"/> 45	<input type="text"/> 3/1/1999	<input type="text"/> No	<input type="text"/> 2.00

Liver Function

Specimen Date	Value	Date	Specimen Date	Specimen Value	Explanations
<input type="text"/> 3/1/1999	<input type="text"/> 49	<input type="text"/> 45	<input type="text"/> 3/1/1999	<input type="text"/> No	<input type="text"/> 19.2

Blood Coagulation

Specimen Date	Value	Date	Specimen Date	Specimen Value	Explanations
<input type="text"/> 3/1/1999	<input type="text"/> 49	<input type="text"/> 45	<input type="text"/> 3/1/1999	<input type="text"/> No	<input type="text"/> 2.00

Urine

Specimen Date	Value	Date	Specimen Date	Specimen Value	Explanations
<input type="text"/> 3/1/1999	<input type="text"/> 49	<input type="text"/> 45	<input type="text"/> 3/1/1999	<input type="text"/> No	<input type="text"/> 19.2

TPMS F

How To

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TPMS

F16.10A

TPMS Patient

Hazardous Drugs: None in Treatment

Virologic Current Therapy: 127 101 IV

Therapeutic Options (127-101)		127		101		127 + 101 Therapies		127		101		127 + 101 Therapies		
Protocol	Drug	Dose	Route	Dose	Route	Dose	Route	Dose	Route	Dose	Route	Dose	Route	
△ ddI, d4T, NFV	2	2	ddI Renal dos. Adj.	d4T Renal dos. adj	q8h	15	\$30.38	✓ AZT (Retrovir/zidovudine)	q12h	18	\$34.06	✓ AZT (Retrovir/zidovudine)	q12h	
△ ddI, d4T, RTV	4	4	ddI Renal dos. Adj.	d4T Renal dos. adj	q12h	9	\$44.32	ddI (Viread/didanosine)	q12h	9	\$44.32	ddC (Hivid/zalcitabine)	q12h	
△ NVP, ABC, EFV	5	5	NVP Renal dos. Adj.	EFV+Renal Dyst	q8h	19	\$43.21	✓ 3TC (Epivir/lamivudine)	q8h	16	\$54.40	✓ d4T (Zefit/stavudine)	q8h	
△ DLV, ABC, EFV	5	5	EFV+Renal Dyst	5	EFV+Renal Dyst	q8h	17	\$46.41	ABC (Zagren/abacavir)	q8h	17	\$46.41	abc	q8h
△ NFV, ABC, EFV	5	5	NVP Renal dos. Adj.	EFV+Renal Dyst	q8h	17	\$46.41	abc	q8h	17	\$46.41	abc	q8h	
△ NVP, NFV, EFV	5	5	NVP Renal dos. Adj.	EFV+Renal Dyst	q8h	17	\$46.41	abc	q8h	17	\$46.41	abc	q8h	

See Note Endoscopy Evaluation

Therapeutic Options (127-101)		127		101		127 + 101 Therapies		127		101		127 + 101 Therapies	
Protocol	Drug	Dose	Route	Dose	Route	Dose	Route	Dose	Route	Dose	Route	Dose	Route
△ tenofovir	22	3TC	IV	101	IV	✓	101	101	IV	101	IV	✓	101
valganciclovir													



- AZT: Medical Condition Alert: This patient has a history of anemia. Use Retrovir with caution due to risk of hematologic toxicity. More Info 171

FiltRankC, Commentary 171

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Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- Epivir 150mg q24h (1 pill/day, \$3.84/day)
- Crixivan 800mg q8h (6 pills/day, \$15.00/day)

(♦ indicates adjusted dosage)

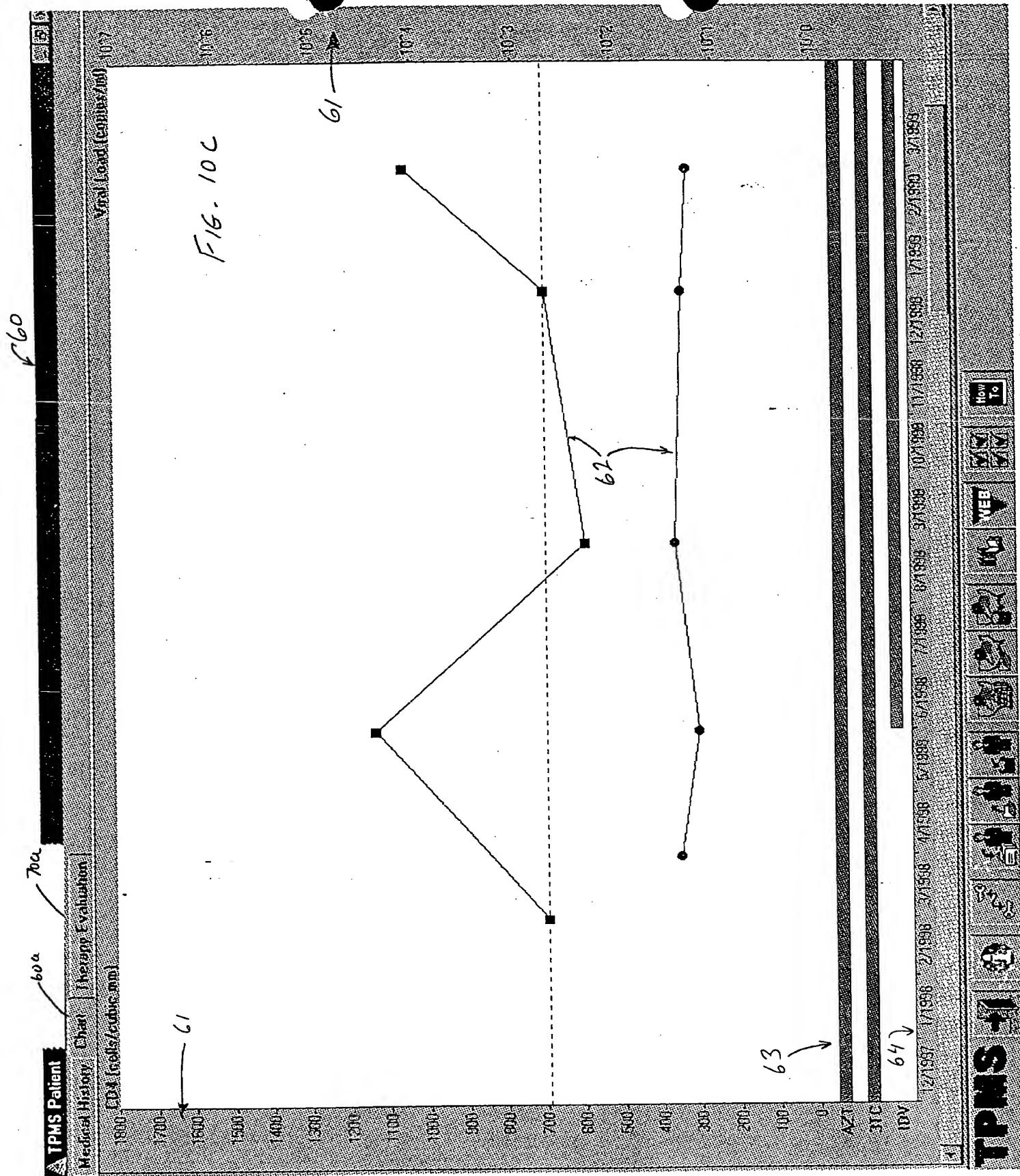
Warning - Resistance Notices

- Resistance Advisory: Retrovir and Epivir ranked lower (+) due to historical virological failure. More Info 364 FiltResF13, Commentary 264

• Dolutegravir: Previous exposure to Dolutegravir but no outcome data. Patients ranked down to More Info 251

TPMS

How To



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Medical History Chart - Headline Evaluation	
Patient ID	demo1
Patient Name	Birth Day: 1/1/1960
Gender	Male
HIV and Viral Load	
HIV (estimated viral load)	37
Current Viral Load	37
Previous Viral Load	37
HIV Genotype	
Phenotype	
Allergy/Hyper	
Intolerance	
Hemoglobin	
Specimen Date	3/1/1999
Value	12
Neutrophils	
Specimen Date	3/1/1999
Value	150
Hepatic Function	
Specimen Date	3/1/1999
Value	49

Boundary and Prequalification Messages

• Poor Viral Suppression Δ : The patient's viral load count either did not decrease $\geq 5 \log$ from the last point or is not below the viral load reduction goal. Unless lab error is at fault, consider changing therapy. More Info FQ1_PeQualA6, Commentary#45

Data Needed Soon - Caution

- No Baseline Viral Load Value: Please specify which viral load value or values (an average of two points) you wish to be set as the baseline viral load value for this patient.

BoundZY, Commentary#41a

MBI

FIG. 10D

TPMS 4

How
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TPMS Patient #60a

Initial Evaluation

General		<input checked="" type="checkbox"/> Run	<input checked="" type="checkbox"/> Computer Print	Date	Value
Patient #	Birth Date	TPMS Number	Weight (kg)	2/1/1999	73.00
ARV naïve		Run	Send Dose	2/1/1999	Yes
Female	Male	Print	Print		
CD4 and Viral Load		Specimen Date	Specimen Value	AIDS Diagnostic	
CD4 (cells/cmm ³)	2/20/1999	350	1/20/1999	175	
Concentrated Load	2/20/1999	31000	V. UNK	0.1	
Previous Viral Load	2/20/1998	19000	V. UNK	0.1	
HIV Serotype		Specimen Date	Specimen Value	Non ARV Drugs	
Phenotype				10/5/1998	
Allergy/Hyper				Bacitracin DS Tablets	oral
Intolerance					12/6/1998
Hemoglobin		Specimen Date	Specimen Value	Serum Standard	
Phenotype	2/1/1999	12.50			
Neurofibrosis		Specimen Date	Specimen Value		
Specimen Date	2/1/1999	1350	2/1/1999	No	
Pancreatitis		Specimen Date	Specimen Value		
Specimen Date	2/1/1999	12.50	2/1/1999	No	
Hepatic Function		Specimen Date	Specimen Value	Renal Function	
Specimen Date	AST (SGOT) U/L	35	2/1/1999	No	2/1/1999
2/1/1999	35		2/1/1999	1.00	1.00
				How Tc	
				Wt	ECG
				HR	ECG
				BP	ECG
				RR	ECG
				SpO ₂	ECG
				Tc	ECG

FIG. 11A

TPMS

Tc

Wt

ECG

HR

BP

RR

SpO₂

70a

Medical History / Clinical Therapy Evaluation

General

Patient ID	Birth Date	TEMS Number
151968		

Gender

Male

Boundary and Prequalification Messages

 UK
 USA

Please be aware that the following boundary and prequalification messages currently apply to the patient.

HIV Genotype

Phenotype

Allergy/Hyper

Tolerance

Hemoglobin

Screen Date	V
2/1/1999	12

Neutrophilic

Screen Date	V
2/1/1999	13

Hepatic Function

Screen Date	V
2/1/1999	35

- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E/ml bDNA) or CD4 counts less than 500 cells/ml (Ann. Int. Med., 1998). PreQualM, Commentary§1

- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PreQualM, Commentary§6

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To

TPMS Patient

Written History: Frank - HIV-Infected Patient

Initial Treatment		Treatment Regimen		Treatment Regimen	
Therapy	Regimen	Therapy	Regimen	Therapy	Regimen
AZT, ddI, 3TC, SQV, SGC	1	1	1	q8h	\$43.46
ddI, 3TC, NFV	1	1	1	q8h	\$34.78
AZT, 3TC, IDV	1	1	1	q8h	\$32.24
AZT, 3TC, NFV	1	1	1	q8h	\$35.81
d4T, 3TC, IDV	1	1	1	q8h	\$31.20
AZT, ddI, RTV, DLV	2	2	DLV+RTV	q8h	\$45.99
ddI, d4T, IDV, NVP	2	2		q8h	\$42.55
d4T, 3TC, RTV	2	2		q12h	\$38.46
AZT, ddI, RTV, NVP	2	2		q12h	\$47.10
See Note	See All	See All	See All	See All	See All

Initial Treatment		Treatment Regimen		Treatment Regimen	
Therapy	Regimen	Therapy	Regimen	Therapy	Regimen
AZT, 3TC, IDV	1	1	1	q8h	\$43.46
ddI, 3TC, NFV	1	1	1	q8h	\$34.78
AZT, 3TC, IDV	1	1	1	q8h	\$32.24
AZT, 3TC, NFV	1	1	1	q8h	\$35.81
d4T, 3TC, IDV	1	1	1	q8h	\$31.20
AZT, ddI, RTV, DLV	2	2	DLV+RTV	q8h	\$45.99
ddI, d4T, IDV, NVP	2	2		q8h	\$42.55
d4T, 3TC, RTV	2	2		q12h	\$38.46
AZT, ddI, RTV, NVP	2	2		q12h	\$47.10
See Note	See All	See All	See All	See All	See All

Initial Treatment		Treatment Regimen		Treatment Regimen	
Therapy	Regimen	Therapy	Regimen	Therapy	Regimen
AZT, 3TC, IDV	1	1	1	q8h	\$43.46
ddI, 3TC, NFV	1	1	1	q8h	\$34.78
AZT, 3TC, IDV	1	1	1	q8h	\$32.24
AZT, 3TC, NFV	1	1	1	q8h	\$35.81
d4T, 3TC, IDV	1	1	1	q8h	\$31.20
AZT, ddI, RTV, DLV	2	2	DLV+RTV	q8h	\$45.99
ddI, d4T, IDV, NVP	2	2		q8h	\$42.55
d4T, 3TC, RTV	2	2		q12h	\$38.46
AZT, ddI, RTV, NVP	2	2		q12h	\$47.10
See Note	See All	See All	See All	See All	See All

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- **WARNING:** Before initiating any antiRetroviral treatment regimen, the complete product information for each therapeutic component should be consulted.
- **Viral Load Testing Required:** Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary 55

F16- 11 C

A2

- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 Eq/ml bDNA) or CD4 counts less than 500 cells/ml. (Ann. Int. Med., 1998). PeQualM, Commentary 61
- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PeQualM, Commentary 66

A3

TPMS

How To

TPMS Patient

Helen Hwang Chait Therapy Evaluation

Therapy: R-TG / ZDV / NRTV / D4T

Evaluated

Use in Orientations

Show Examples

Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- Virex 200mg q12h (4 pills/day, \$6.78/day)
- Norvir 600mg q12h (12 pills/day, \$22.26/day)
- Rescriptor 400mg q8h (12 pills/day, \$7.39/day)

F16. 11D

- AZT: Interrupt Retroviruse if anemia and/or neutropenia develops. More Info 036 DosGenA, Commentary³⁶
- ddI: When treatment with other drugs known to cause pancreatic toxicity is required (for example, IV pentamidine), suspension of Videx should be considered. CmtGenA, Commentary¹³
- ddI: If patients develop symptoms of neuropathy, Videx therapy should be interrupted. DosGenB, Commentary⁴⁰
- ddI: Clinical signs suggestive of pancreatitis should prompt dose suspension of Videx and careful evaluation of the possibility of pancreatitis. Only after pancreatitis has been ruled out should dosing be resumed. DosGenB, Commentary³⁹
- DLV: Skin rash attributable to Rescriptor may occur during first 21 days. More Info 054 CmtGenS, Commentary⁵⁴
- ddI: Videx should not be administered with a prescription antibiotic containing any form of tetracycline. CmtGenA, Commentary⁵
- ddI: Plasma concentrations of some quinolone antibiotics are decreased when administered with antacids containing magnesium or aluminum. Therefore, doses of quinolone antibiotics should not be administered within 2 hours of taking Videx. CmtGenA, Commentary¹⁶
- RTV: Monitor for decreased AUC of RTV and associated adverse events when concomitant with use of drugs that increase CYP3A activity (including tobacco). More Info 026 CmtGenH, Commentary²⁶

TPMS

How To

WEB

PDF

TPMS Patient

Medical History | Chemotherapy Evaluation

<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old
<input type="checkbox"/> AZT, ddI, 3TC, SQV, SGC d4T, 3TC, NFV	<input checked="" type="checkbox"/> Show Abstract for Retrovir	<input type="checkbox"/> Show Abstract for Raltegravir
<input type="checkbox"/> AZT, 3TC, IDV AZT, 3TC, NFV	<input type="checkbox"/> Show Abstract for Epivir	<input type="checkbox"/> Show Abstract for Emtricitabine
<input type="checkbox"/> d4T, 3TC, IDV	<input type="checkbox"/> Show Abstract for Stavudine	<input type="checkbox"/> Show Abstract for Zalcitabine
<input type="checkbox"/> AZT, ddI, RTV, DLV ddI, d4T, IDV, NVP	<input type="checkbox"/> Show Abstract for Zidovudine	<input type="checkbox"/> d4T (Zidovudine)
<input type="checkbox"/> d4T, 3TC, RTV AZT, ddI, RTV, NYV	<input type="checkbox"/> Show Abstract for Zidovudine	<input type="checkbox"/> ABC (Zagren / abacavir)
<input type="checkbox"/> See More	<input type="checkbox"/> Print Top 10 Therapeutic Options	<input type="checkbox"/> Print Top 10 Therapeutic Options
<input type="checkbox"/> Print All Therapeutic Options	<input type="checkbox"/> Print Summary	<input type="checkbox"/> Print Summary
<input type="checkbox"/> Print Top Therapeutic Options	<input type="checkbox"/> Print Summary	<input type="checkbox"/> Print Summary

Viral Load Testing		Viral Load Testing		Viral Load Testing	
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<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old
<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old
<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old	<input type="checkbox"/> Hide Column: New [+] Hide Column: Old

<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape
<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape
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<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape
<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape	<input type="checkbox"/> Show Long Therape [+] Show Short Therape

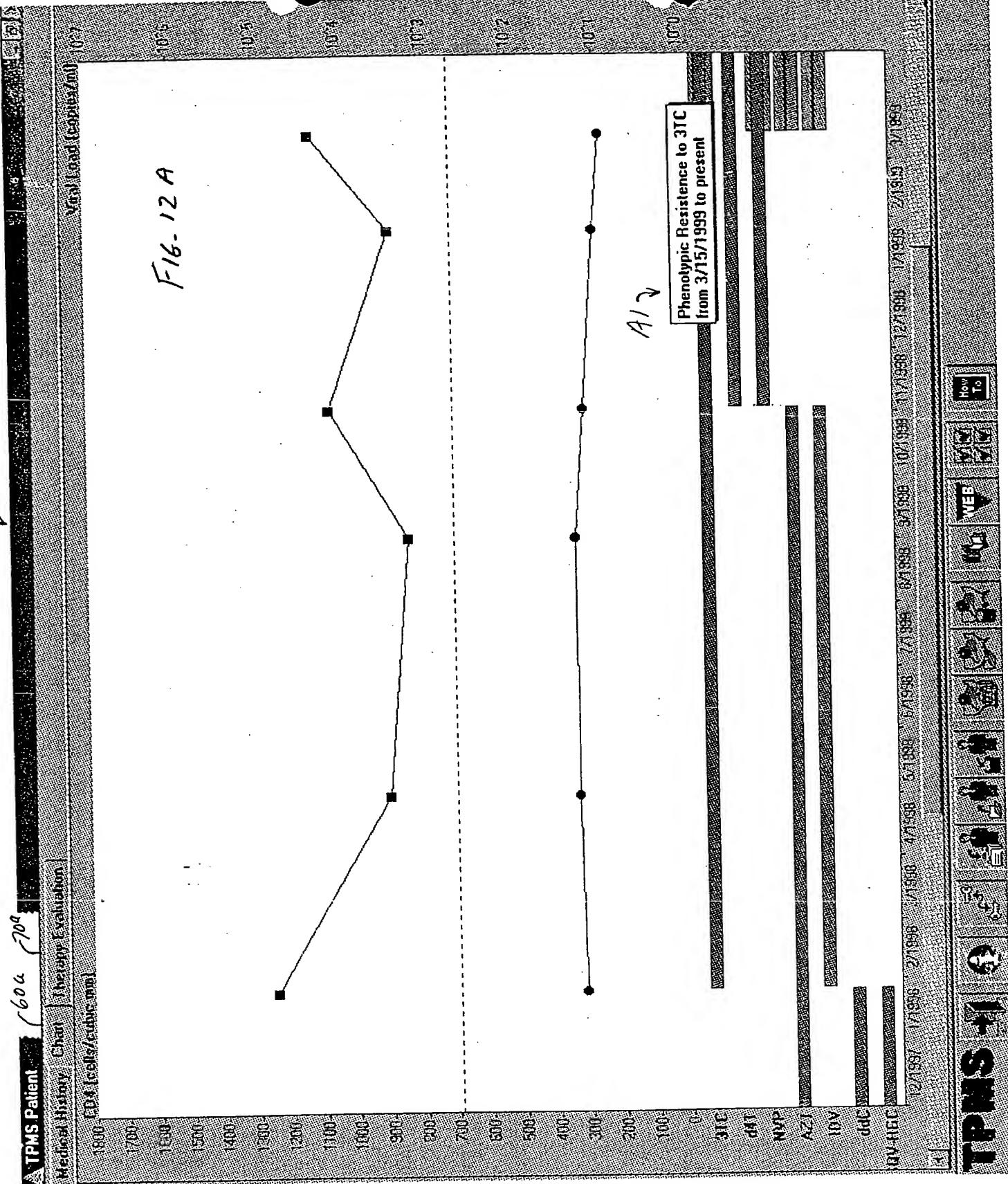
F16-11E

- Therapy Initiation: Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E/g/ml tDNA) or CD4 counts less than 500 cells/ μ l (Ann. Int. Med., 1998). PaQualM, Commentary6
- Combination Therapy Recommended: Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PaQualM, Commentary6

TPMS

How To

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TPMS PatientPatient Name: Chris Last Name: UnknownEvidential Genotype: 3TC/NVPLast Genotype: 3TC/NVPEvaluated: NeverLast Evaluated: NeverLast Rejection: NeverLast Rejection Reason: NeverLast Rejection Date: NeverLast Rejection ID: NeverLast Rejection Result: NeverLast Rejection Comment: NeverLast Rejection Status: NeverLast Rejection Status Date: NeverLast Rejection Status Comment: NeverLast Rejection Status Result: NeverLast Rejection Status Comment: NeverTPMS ID: 42

How To

Therapy		EfV	ddI	ddC	d4T	Rifabutin	EpV	ADV	ddI
<input type="checkbox"/> ddI, d4T, NVP	2	2				Rifabutin+NVP	q8h	15	\$33.88
<input checked="" type="checkbox"/> ddI, d4T, EFV	5	5				Rifabutin	q12h	9	\$28.44
<input type="checkbox"/> ddI, NVP, EFV	5	5				Rifabutin+NVP	q8h	16	\$38.50
<input type="checkbox"/> ddI, NVP, EFV	5	5				Rifabutin+NVP	q8h	14	\$40.24
<input type="checkbox"/> d4T, NVP, EFV	5	5				Rifabutin+NVP	q8h	15	\$38.77
<input type="checkbox"/> ddC, NVP, EFV	5	5				Rifabutin+NVP	q8h	8	\$28.71
<input checked="" type="checkbox"/> ddC, d4T, EFV	5	5				Rifabutin+NVP	q8h	8	\$28.71
<input type="checkbox"/> See Note	1	1				Rifabutin+NVP	q8h	8	\$28.71

Reason Being Evaluated:
Rejected

III THERAPY REJECTED !!!

This therapy was rejected for the following reason(s) Additional information about the therapy is provided but this therapy is NOT advisable

- Viramune (nevirapine/NVP) Resistance Advisory: According to the last genotype data entered, the patient's virus currently has mutation(s) which is/are associated with resistance to Viramune. FillMutE, RejectionS4

- Resistance Advisory: According to the last genotype data entered, the patient's virus currently has the following mutations; M184V [RT]. The genotype test displays evidence of the M184V/M184I mutation which is associated with resistance to 3TC. However, this mutant has increased sensitivity to the anti-retroviral activity of AZT and ADV so an AZT/3TC or AZT/ADV combination is still useable. Therefore combinations which contain AZT/3TC and AZT/ADV are shown as therapy options although these therapies have been ranked down +5 in favor of three drug combinations with no resistant mutants. FillMutB, RejectionS1

- Epivir and Viramune Resistance Advisory: The patient's last phenotypic assay demonstrates phenotypic resistance to Epivir and Viramune, therefore, therapies containing Epivir and Viramune are not recommended at this time. FillResC, RejectionS2

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CAUTION

YELLOW ALERT

✓✓3

- NVP Δ: Drug Interaction Alert: Patient is currently taking rifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs

TPMS

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Medical History | Chief | Therapy Evaluation

General		<input checked="" type="checkbox"/> H	<input checked="" type="checkbox"/> Enzyme	<input checked="" type="checkbox"/> Comment	Populations	Date	Value
Patient Features	Birth Date	1/1/1950	YFMN Number		Weight (kg)	1/28/1999	60.00
Phenotype	Gender	Male		Sex	Sex/drug	1/28/1999	Yes
ID 4 and Viral Load		Specimen Date	Value	Specimen Date	Value	Specimen Date	Value
CNA	3/15/1999	240	1/28/1999	45	1/28/1999	10	
Ceftriaxone	3/15/1999	21500	D/ml		V/ml		
Current Viral load	3/15/1999	3600	V/ml		V/ml		
Previous Viral load							
HIV Genotype		Specimen Date	Value	Specimen Date	Value	Specimen Date	Value
L101[P]	3/15/1999	M461[P], M461[R1], Y181					
<p>• NVPΔ: Drug Interaction Alert: Patient is currently taking rifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs should only be used in combination if clearly indicated and with careful monitoring. CmDIP, Commodity³³</p>							
Hemoglobin		Specimen Date	Value (g/dL)	Specimen Date	Value	Specimen Date	Value
		1/28/1999	15.00		1/28/1999	No	
Neutrophils		Specimen Date	Value (x10 ⁹ /mm ³)	Specimen Date	Value	Specimen Date	Value
		1/28/1999	1500		1/28/1999	No	
Hepatic Function		Specimen Date	AST (U/L)	Specimen Date	LDH (U/L)	Specimen Date	Gamma Gt (U/L)
		1/28/1999	25		25	1/28/1999	1.00
<p>• V3</p> <p>5/1999</p> <p>5/1999</p>							

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TPMS-1

How
To